

## Male contraceptive formulation comprising Norethisterone

### FIELD OF INVENTION

a1 <sup>5</sup> The invention relates to a formulation of a male contraceptive comprising a progestin as well as methods of male contraception utilising progestins. Moreover, the invention relates to formulation further comprising an androgen as well as to methods using formulations comprising a progestin and an androgen so as to suppress  
10 spermatogenesis.

### GENERAL BACKGROUND

Contraceptive methods for men are considered an essential component of world-wide  
15 reproductive health (Nieschlag and Behre; *Testosterone: action, deficiency, substitution*, 1998, Springer, Berlin, p 514). Hormonal methods of male contraception offer the advantages of high-reversibility and efficacy. In hormonal male contraception, the suppression of spermatogenesis is sought through the suppression of the gonadotropins leuteinizing hormone (LH) and follicle stimulating hormone (FSH) to undetectable levels  
20 within the endocrine feedback mechanism operating between the pituitary gland and the hypothalamus. Disadvantageously, suppression of these gonadotropins also induces symptoms related to androgen deficiency (Nieschlag and Behre; 1998, pp 513-528).

Male contraceptive methods seek to suppress FSH and LH, resulting in a depletion of  
25 intratesticular testosterone and cessation of spermatogenesis, whilst substituting peripheral testosterone with another androgen. This androgen has typically been testosterone itself and serves the endocrine androgenic role of testosterone such as to maintain libido, male sex characteristics, protein anabolism, hematopoiesis and others. In short, the objective is to deplete the testes of testosterone whilst maintaining levels in  
30 the general circulation.

The suppression of spermatogenesis by administration of testosterone alone have been ineffective in inducing azoospermia. The administration of gonadotropin releasing hormone antagonists (GnRH antagonists) has circumvented part of the problem with the

administration of high levels of testosterone alone but GnRH antagonists are unattractive for clinical use in their current preparations and are generally expensive to prepare.

The use of either the progestins cyproterone acetate or levonorgestrel were either  
 5 ineffective in the suppression of spermatogenesis or, in higher doses, led to significant decreases in red blood cell count (Merrigiola *et al*, 1998; Merrigiola *et al*, 1997; Merrigiola *et al*, 1996; Bebb *et al*, 1996).

The use of a mixture of two compounds, one an androgen and another an estrogen, in  
 10 combination is disclosed in US 4,210,644.

A method towards the inhibition of spermatogenesis in men by administering testosterone percutaneously or orally and the progestin norethisterone acetate orally has been disclosed (Guerin and Rollet, 1988). However, high doses relative to amounts required by  
 15 the present invention of each component were required to achieve azoospermia.

Thus, effective and efficient methods of male contraceptive by use of a male contraceptive formulation are not currently available.

## 20 BRIEF DESCRIPTION OF THE INVENTION

The invention disclosed herein solves this problem by providing

in a first aspect, a male contraceptive formulation comprising an effective amount of a  
 25 progestin, wherein said progestin possesses both androgenic and estrogenic properties. Most preferably, the formulation comprises the progestin norethisterone enanthate. Furthermore, the formulation may comprise norethisterone (NET), or derivatives thereof, and an androgen. Most preferably, in formulations comprising NET, or derivatives thereof and an androgen, the androgen is testosterone undecanoate.

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A second aspect of the invention relates to the use of NET, or derivatives thereof, for the preparation of a pharmaceutical composition for use as a male contraceptive. Particularly, the use of NET enanthate or NET acetate for the preparation of a pharmaceutical composition for use as a male contraceptive is defined herein.

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Similarly, the use of a combination of NET, or derivatives thereof, and an androgen for the preparation of a pharmaceutical composition for use as a male contraceptive is defined herein. Furthermore, the use of a combination of NET enanthate or NET acetate and testosterone undecanoate for the preparation of a pharmaceutical composition for use as

5 a male contraceptive.

The use of a combination of NET esters and testosterone, or derivatives thereof, for the preparation of a male contraceptive formulation is also defined herein.

- 10 A third aspect of the invention relates to a method of providing male contraception comprising administering to an individual NET or a derivative thereof in an amount sufficient to suppress spermatogenesis.

#### DETAILED DESCRIPTION OF THE INVENTION

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In the present application, a number of terms are used which are commonly used in the pharmaceutical field. An explanation of some of the special terms and concepts relevant to the present is given in the following items:

20 *estrogenic*

-The term *estrogenic* when used alone or when applied to the term *properties* is intended to imply the characteristics of compounds exhibiting estrogen-like activities as defined by *in vitro* receptor-binding assays, transactivation assays, or *in vivo* assays as described in Example 2.

25

*androgenic*

-The term *androgenic* when used alone or when applied to the term *properties* is intended to imply the characteristics of compounds exhibiting androgen-like activities as defined by *in vitro* antiproliferation assays, transfection assays, or *in vivo*

30 assays as described in Example 3.

*spermatogenesis*

- The term *spermatogenesis* is intended to mean the overall process of gametogenesis in the male. Spermatogenesis takes place in the seminiferous tubule and

is directly regulated by levels of follicle stimulating hormone and androgen at the periphery of the seminiferous tubule, particularly upon the Sertoli cells.

#### *azoospermia*

- 5                   - The term *azoospermia* represents a sperm content in semen sample below 1 million per mL (mill/mL) approaching levels of zero sperm content and are the result of suppression of spermatogenesis

#### *oligozoospermia*

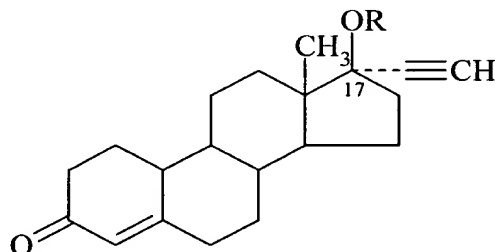
- 10                  - The term *oligozoospermia* represents a sperm content in semen between 20 and one million per mL (mill/mL) and are the result of inhibited levels of spermatogenesis.

#### *Pearl-Index*

- 15                  - The term *Pearl Index* is intended to indicate the number of pregnancies per 1200 months.

#### *NET derivatives*

- 20                  -The term *derivatives* when applied to the term *NET* is intended to mean derivatives of 19-norethisterone, known under a variety of names including 17-Hydroxy-19-norpreg-4-en-20-yn-3-one. NET derivatives are intended to mean any modification of the 17-hydroxyl group without changing the oxidation state of carbon 17 of the compound as depicted in formula A. Thus, R can be selected from any number of chemical groups. These include groups so as to form ether, carboxylic ester, silyl, phosphate, sulphonate  
25 and sulfate.



A

The term ether is intended to indicate that R is selected from the group comprising of optionally substituted C<sub>1-18</sub>-alkyl, C<sub>2-18</sub>-alkenyl, C<sub>2-18</sub>-alkynyl, C<sub>3-8</sub>-cycloalkyl, heterocyclyl,

- and aryl. In the present context the term " $C_{1-18}$ -alkyl" used alone or as part of another group designates a linear or branched saturated hydrocarbon group having from one to eighteen carbon atoms, such as methyl, ethyl, n-propyl, isopropyl, cyclopropyl, n-butyl, isobutyl, sec-butyl, tert-butyl, cyclobutyl, pentyl, isopentyl, neopentyl, cyclopentyl, hexyl, cyclohexyl, pivalyl, undecanyl, dodecanyl, myristyl (tetradecanyl), palmityl (hexadecanyl) and stearyl (octadecanyl). In the present context the term " $C_{2-18}$ -alkenyl" is intended to mean a linear or branched hydrocarbon group having from two to eighteen carbon atoms and containing one or more double bonds. Examples of  $C_{2-18}$ -alkenyl groups include olefins such as allyl, homo-allyl, vinyl, crotyl, butenyl, pentenyl, hexenyl, heptenyl and octenyl. Examples of  $C_{2-8}$ -alkenyl groups with more than one double bond include butadienyl, pentadienyl, hexadienyl, heptadienyl, hexatrienyl, heptatrienyl and octatrienyl groups as well as branched forms of these. In the present context the term " $C_{2-18}$ -alkynyl" is intended to mean linear or branched hydrocarbon groups containing from two to eighteen carbon atoms and containing one or more triple bonds. Examples of  $C_{2-18}$ -alkynyl groups include acetylene, propynyl, butynyl, pentynyl, hexynyl, heptynyl and octynyl groups as well as branched forms of these. In the present context the term " $C_{3-8}$ -cycloalkyl" is intended to cover three-, four-, five-, six-, seven-, and eight-membered rings comprising carbon atoms only whereas the term "heterocyclyl" is intended to mean three-, four-, five-, six-, seven-, and eight-membered rings wherein carbon atoms together with from 1 to 3 heteroatoms constitute said ring. The heteroatoms are independently selected from oxygen, sulphur, and nitrogen. Such  $C_{3-8}$ -cycloalkyl and term heterocyclyl rings may contain no unsaturated bonds or may contain one or more unsaturated bonds, however, if present, situated in such a way that an aromatic  $\pi$ -electron system does not arise.
- Examples of " $C_{3-8}$ -cycloalkyls" are cyclopropane, cyclobutane, cyclopentane, cyclopentene, cyclopentadiene, cyclohexane, cyclohexene, 1,3-cyclohexadiene, 1,4-cyclohexadiene, cycloheptane, cycloheptene, 1,2-cycloheptadiene, 1,3-cycloheptadiene, 1,4-cycloheptadiene and 1,3,5 cycloheptatriene.
- Examples of "heterocyclyls" are 2*H*-thiopyran, 3*H*-thiopyran, 4*H*-thiopyran, tetrahydrothiopyran, 2*H*-pyran, 4*H*-pyran, tetrahydropyran, piperidine, 1,2-dithiin, 1,2-dithiane, 1,3-dithiin, 1,3-dithiane, 1,4-dithiin, 1,4-dithiane, 1,2-dioxin, 1,2-dioxane, 1,3-dioxin, 1,3-dioxane, 1,4-dioxin, 1,4-dioxane, piperazine, 1,2-oxathiin, 1,2-oxathiane, 4*H*-1,3-oxathiin, 1,3-oxathiane, 1,4-oxathiin, 1,4-oxathiane, 2*H*-1,2-thiazine, tetrahydro-1,2-thiazine, 2*H*-1,3-thiazine, 4*H*-1,3-thiazine, 5,6-dihydro-4*H*-thiazine, 4*H*-1,4-thiazine,

tetrahydro-1,4-thiazine, 2*H*-1,2-oxazine, 4*H*-1,2-oxazine, 6*H*-1,2-oxazine, 2*H*-1,3-oxazine, 4*H*-1,3-oxazine, 4*H*-1,4-oxazine, morpholine, trioxane, 4*H*-1,2,3-trithiin, 1,2,3-trithiane, 1,3,5-trithiane, hexahydro-1,3,5-triazine, tetrahydrothiophene, tetrahydrofuran, pyrroline, pyrrolidine, pyrrolidone, pyrrolidione, pyrazoline, pyrazolidine, imidazoline, imidazolidine, 5 1,2-dioxole, 1,2-dioxolane, 1,3-dioxole, 1,3-dioxolane, 3*H*-1,2-dithiole, 1,2-dithiolane, 1,3-dithiole, 1,3-dithiolane, isoxazoline, isoxazolidine, oxazoline, oxazolidine, thiazoline, thiozolidine, 3*H*-1,2-oxathiole, 1,2-oxathiolane, 5*H*-1,2-oxathiole, 1,3-oxathiole, 1,3-oxathiolane, 1,2,3-trithiole, 1,2,3-trithiolane, 1,2,4-trithiolane, 1,2,3-trioxole, 1,2,3-trioxolane, 1,2,4-trioxolane, 1,2,3-triazoline and 1,2,3-triazolidine.

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The term "aryl" is intended to comprise heteroaryl and fused aryl systems. When used in themselves or when described as substituted or optionally substituted are intended to mean the class of cyclic compounds in which themselves or moieties thereof possess chemical aromaticity. Aryls comprise of radicals of optionally mono-, di-, tri-, or tetra- 15 substituted phenyl. In the present context the term "aryl" used alone or as part of another group is intended to mean an aromatic system. The term "heteroaryl" is intended to mean an aryl group where one or more carbon atoms have been replaced with heteroatoms such as nitrogen, sulphur, and/or oxygen atoms. These include a monoradical selected from the group comprising an optionally substituted, such as mono-, di-, tri-, or tetra- 20 substituted furanyl, thiophenyl, pyrrolyl, phenoxazonyl, oxazolyl, thiazolyl, isothiazolyl, imidazolyl, pyrazolyl, isoxazolyl, imidazolyl isothiazolyl, oxadiazolyl, furazanyl, triazolyl, thiadiazolyl, piperidinyl, pyridinyl, pyridazinyl, pyrimidinyl, pyrazinyl, pyrazolyl and triazinyl. "Fused aryls" generally comprise fused ring systems between at least two aryls, at least one aryl fused with at least one heteroaryl, two heteroaryls, at least one aryl fused with a 25 C<sub>3-8</sub>-cycloalkyl, at least one aryl fused with a heterocyclyl, at least one heteroaryl fused with a heterocyclyl or at least one heteroaryl fused with a C<sub>3-8</sub>-cycloalkyl. Fused ring systems, be it between aromatic systems, or between an aromatic system and a non-aromatic ring system are intended to mean where at least two rings share at least one chemical bond. Examples of fused aryls comprise naphthalenyl, phenanthrenyl, 30 anthracenyl, acenaphthylenyl, fluorenyl, indenyl, indolyl, isoindolyl, indolinyl, benzofuranyl, benzothiophenyl, benzopyrazolyl, indazolyl, benzimidazolyl, benzthiazolyl, purinyl, quinoliziny, quinolinyl, isoquinolinyl, cinnolinyl, coumaranyl, coumarinyl, phthalazinyl, quinazolinyl, quinoxalinyl, naphthyridinyl, pteridinyl, chromanyl, isochromanyl, thienofuranyl, carbazolyl, acridinyl, phenazinyl, phenothiazinyl, 35 phenoxazinyl, thianthrenyl and azulenyl.

The term *carboxylic ester* when applied to *NET* is intended to mean the carboxylic acid derivatives of *NET* wherein *R* is the monoradical  $-C(=O)-R'$ . *R'* can be optionally substituted  $C_{1-18}$ -alkyl,  $C_{2-18}$ -alkenyl,  $C_{2-18}$ -alkynyl,  $C_{3-8}$ -cycloalkyl, heterocyclyl, and aryl as defined *supra*. Specific examples of carboxylic acid derivatives of *NET* are wherein the carboxylic acid or derivative of carboxylic acid used in the condensation to *NET* are selected from optionally substituted acetic acid, propionic acid, butanoic acid, pentanoic acid, hexanoic acid, heptanoic acid, octanoic acid (to give the enanthate), undecanoic acid, benzoic acid, toluic acid, salicylic acid, 3-(cyclopentyl)-propionic acid (to give the cypionate), cyclohexanecarboxylic acid, and 4-(butyl)-cyclohexanecarboxylic acid (to give the bucilate), pivalic acid, lauric acid, myristic acid, palmitic acid, stearic acid, acrylic acid, oleic acid and nicotinic acid.

The terms silyl, sulphonate and sulfate are intended to indicate that *R* is the radicals of  $-SiR'R''R'''$ ,  $-SO-R'$ ,  $-SO_2-R'$ , respectively, wherein *R'*, *R''* and *R'''* are independently selected from the group  $C_{1-18}$ -alkyl,  $C_{2-18}$ -alkenyl,  $C_{2-18}$ -alkynyl,  $C_{3-8}$ -cycloalkyl, heterocyclyl, and aryl as defined *supra*. The term phosphate is intended to mean indicate that *R* is the radicals of  $-P(O)(OR')(OR'')(OR''')$ , wherein wherein *R'*, *R''* and *R'''* are independently selected from the group  $C_{1-18}$ -alkyl,  $C_{2-18}$ -alkenyl,  $C_{2-18}$ -alkynyl,  $C_{3-8}$ -cycloalkyl, heterocyclyl, and aryl as defined *supra* or hydrogen.

### *Progestin*

-The term progestin is synonymous with the term progestagen, and comprises a class of hormones naturally present in the body as well as synthetic and semi-synthetic derivatives thereof.

A unique approach of the contraceptive formulation disclosed herein is that of an effective amount of a single progestin possessing both androgenic and estrogenic properties.

In a male contraceptive formulation, the use of a compound with strong androgenic activity suppresses testicular spermatogenesis, and minimises the potential of side effects associated with androgen deficiency whilst feedback regulating LH and FSH from the hypothalamus and pituitary gland. The known estrogenic effect to suppress gonadotropin secretion results in a surprisingly fast and profound suppression of LH and FSH. It has surprisingly been found that the use of a single compound with estrogenic and

androgenic properties is an effective means of suppressing spermatogenesis. A formulation comprising a single compound possessing estrogenic and androgenic properties has been found to be effective in male contraception

- 5 The formulation defined by the invention has both estrogenic and androgenic effects. The progestin norethisterone (NET) possesses both these properties and is therefore a particularly preferred choice of progestin. Equally, norethisterone derivatives are suitable to fulfil the dual role of the progestin. In particular, carboxylic esters of the 17-hydroxy group are especially preferred embodiments of the invention. Norethisterone acetate  
10 (NET-A) and norethisterone enanthate (NET-EN) are considered to be particularly attractive embodiments.

In a particularly interesting embodiment, the formulation further comprises an androgen. Testosterone or testosterone derivatives are preferred such androgens. Most preferably,  
15 testosterone esters are selected as androgens. As is known to the person skilled in the art, a whole host of testosterone esters are effective androgens and may be selected from the group comprising testosterone propionate, testosterone undecanoate, testosterone enanthate, testosterone cypionate and testosterone buciclate. In a particularly preferred embodiment, the androgen is testosterone undecanoate.

20

It is preferred that in the formulations comprising a progestin and an androgen, the progestin be selected from NET or its derivatives, such as its carboxylic esters. Particularly, it is preferred that in these instances, the progestin be NET enanthate or NET acetate, most particularly NET enanthate. One such preferred formulation comprises  
25 testosterone undecanoate and norethisterone enanthate.

The use of a compound, such as a NET derivative, having strong androgenic properties and estrogenic properties is advantageous in that this strong androgenic activity minimises the potential of side effects related to androgen deficiency. The estrogenic  
30 activity of NET, on the other hand, suppresses gonadotropin secretion. A formulation comprising a progestin and androgen effectively and rapidly suppresses spermatogenesis in trial studies. Without being limited to a particularly mechanism, it is theorised that the rapidity of the suppression, in comparison to trials using progestins which are not strongly androgenic and estrogenic but instead are anti-androgenic or only weakly androgenic, is  
35 not solely due to the anti-gonadotropin activity of NET, but also due to an additional direct



and strong effect on the testes. This direct testicular effect is likely to be mediated through the androgenic and estrogenic activity of NET.

#### *Doses*

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Preferably, the formulation comprises a dose of NET or of the NET derivative corresponding to a daily release of the NET derivative ranging between 1 and 10 mg, such as 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 mg.

- 10 Correspondingly, the dose of the NET derivative should range between 100 and 500 mg for each 6-week administration, particularly between 200 and 400 mg for each 6-week administration or about 200 mg for each 6-week administration.

The levels of progestin, such as NET or its derivatives, are to be sufficient to suppress spermatogenesis. In general, the levels of progestin are to be sufficient to induce oligozoospermia or azoospermia. In men whose sperm has a less than normal fertilising capacity, either as a result of the contraceptive method, other treatments or naturally, complete azoospermia may not be required for effective contraception. In such cases, levels of progestin such as NET need merely be sufficient to induce oligozoospermia.

20

Accordingly, the formulation may comprise progestin in sufficient amounts to lower the sperm concentration to not more than 3 million/ mL of semen, such as not more than 2 million/ mL, 1 million/ mL, 0,5 million/ mL , 0.25 million/ mL, or 0,1 million/ mL. It is preferred however that the progestin be in sufficient amounts so as to lower the sperm concentration to not more than 0.1 million/ mL.

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Furthermore, the levels of progestin in the formulation may be such that upon following a method disclosed herein, the Pearl-index is not more than 1.4, such as not more than 1.2, 1.0, 0.8, 0.7, 0.6, 0.5, 0.4, 0.3, 0.2, or not more than 0.1.

30

The effective levels of progestin in the circulation should preferably be sustained for not less than 1 week. The formulation and/or consequent method of administration are preferably designed so as to maintain effective levels of progestin in the system for as long as desired by the man, such as for not less than 2, 3, 4, 5, 6, 7,8, 9, 10, 11, or 12 weeks.

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As stated earlier in relation the embodiments of the formulation *not* further comprising an androgen, the formulation comprising a combination of NET or derivatives thereof *and* the androgen must also be in sufficient amounts to suppress spermatogenesis.

5

Preferably, the formulation comprising NET and an androgen comprises a dose of NET or of the NET derivative corresponding to a daily release of the NET derivative ranging between 1 and 10 mg, such as 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 mg. Correspondingly, the dose of the NET derivative should range between 100 and 500 mg for each 6-week  
10 administration, particularly between 200 and 400 mg for each 6-week administration or about 200 mg for each 6-week administration.

In formulations comprising the progestin and an androgen, the dose of the latter should correspond to a daily release of the testosterone derivative corresponding to a daily  
15 release of testosterone in amounts ranging between 5 and 35 mg, such as preferably corresponding to a daily dose of testosterone ranging between 15 and 30 mg.

In such embodiments comprising an androgen and wherein the androgen is testosterone undecanoate, a dose of testosterone undecanoate ranging between 800 and 1500 mg for  
20 each 6-week administration is anticipated, such as a dose of about 1000 mg for each 6-week administration.

In formulations comprising the progestin and an androgen, wherein the progestin is NET enanthate or NET acetate, the dose of the latter should correspond to a daily release of  
25 the NET ester ranging between 1 and 10 mg. Moreover, the dose of the NET derivative preferably ranges between 100 and 500 mg for each 6 week administration, such as between 150 and 250 mg for each 6 week administration such as corresponding to dose of NET acetate of about 200 mg for each 6 week administration. In this same formulation, the dose of the androgen may correspond to a daily release of the testosterone derivative  
30 corresponding to a daily release of testosterone in amounts ranging between 5 and 35 mg, such as preferably corresponding to a daily dose of testosterone ranging between 15 and 30 mg. Accordingly, the dose of the testosterone derivative may correspond to a dose of testosterone between 500 and 1200 mg for each 6-week administration, such as corresponding to a dose of testosterone of about 1000, 800, or 650 mg for each 6-week  
35 administration. Preferably, the androgen is testosterone undecanoate.

The formulation comprising NET or derivatives thereof and the androgen should preferably be in sufficient amounts to lower the sperm concentration to not more than 3 million/ mL, most preferably to sperm concentrations of not more than 0.1 million/ mL.

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It is an objective of the invention to provide a formulation comprising NET or derivatives thereof and the androgen in sufficient amounts such that Pearl-index is lowered to not more than 1.4, such as not more than 1.2, 1.0, 0.8, 0.7, 0.6, 0.5, 0.4, 0.3, 0.2, or not more than 0.1.

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Moreover, a further objective of the invention is to provide a formulation wherein effective circulating levels of the combination of progestin and androgen are sustained for not less than 1 week. Preferably, the formulation and consequent method of administration are such that effective circulating levels of the combination of progestin and androgen are sustained for not less than 2 weeks, such as not less than 4 weeks, 6 weeks, 8 weeks, 10 weeks or 12 weeks.

#### *Methods of Administration*

20 The method of administration should preferably be by non-oral means such as by intramuscular injection, subcutaneous implant, transdermal patch or percutaneous application. Thus, the formulation may be adapted to be administered via intramuscular injection, intravenous injection, subcutaneous implantation, subcutaneous injection or transdermal preparation.

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Intramuscular injections of NET derivatives has led to surprisingly effective serum levels with regards to suppression of spermatogenesis. Given the pharmacodynamics of NET derivatives, this method of administration is considered a particularly attractive embodiment towards achieving effective amounts of the progestin in the blood stream.

30 Moreover, by circumventing the hepatic first pass, lower doses are required to achieve effective amounts.

However, given that the method of administration is a means to an end, that is to say a method of delivering the progestin in the formulation in any of its embodiments at an effective level, other methods of administration are anticipated. The pharmaceutical

other methods \*

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formulation may be administered parenterally by injection, infusion or implantation (intravenous, intramuscular, intraarticular, subcutaneous or the like) in dosage forms, formulations or e.g. suitable delivery devices or implants containing conventional, non-toxic pharmaceutically acceptable carriers and adjuvants.

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The formulation and preparation of such compositions is well-known to those skilled in the art of pharmaceutical formulation. Specific formulations can be found in the textbook entitled "Remington's Pharmaceutical Sciences".

- 10 Compositions for parenteral use may be presented in unit dosage forms, e.g. in ampoules, or in vials containing several doses and in which a suitable preservative may be added. The composition may be in form of a solution, a suspension, an emulsion, an infusion device or a delivery device for implantation or it may be presented as a dry powder to be reconstituted with water or another suitable vehicle before use. One or more
- 15 covehicles, such as ethanol, may also be employed. Apart from the active drug substance, the compositions may comprise suitable parenterally acceptable carriers and/or excipients or the active drug substance may be incorporated into microspheres, microcapsules, nanoparticles, liposomes or the like for controlled release. Furthermore, the composition may, in addition, conveniently comprise suspending, solubilising,
- 20 stabilising, pH-adjusting agents and/or dispersing agents.

- As indicated above, the pharmaceutical compositions according to the invention may comprise the active drug substances in the form of a sterile injection. To prepare such a composition, the suitable active drug substances are dissolved or suspended in a
- 25 parenterally acceptable liquid vehicle. Among acceptable vehicles and solvents that may be employed are water, water adjusted to a suitable pH by addition of an appropriate amount of hydrochloric acid, sodium hydroxide or a suitable buffer, 1,3-butanediol, Ringer's solution and isotonic sodium chloride solution. The aqueous formulation may also contain one or more preservatives, for example, methyl, ethyl or n-propyl p-
  - 30 hydroxybenzoate. In cases where the progestin, the androgen, or both are only sparingly or slightly soluble in water, a dissolution enhancing or solubilising agent can be added or the solvent may apart from water comprise 10-60% w/w of propylene glycol or the like.

- As mentioned, controlled release parenteral compositions may be in form of aqueous
- 35 suspensions, microspheres, microcapsules, magnetic microspheres, oil solutions, oil

suspensions, emulsions or the active drug substance may be incorporated in biocompatible carrier(s), liposomes, nanoparticles, implants or infusion devices.

Materials for use in the preparation of microspheres and/or microcapsules are, e.g.,  
 5 biodegradable/bioerodible polymers such as polyglactin, poly-(isobutyl cyanoacrylate), poly(2-hydroxyethyl-L-glutamine) and poly(lactic acid).

Biocompatible carriers which may be used when formulating a controlled release parenteral formulation are, e.g., carbohydrates such as dextrans, proteins such as  
 10 albumin, lipoproteins or antibodies.

Materials for use in implants are, e.g., non-biodegradable as, e.g., polydimethylsiloxane, or biodegradable such as, e.g., poly(caprolactone), poly(lactic acid), poly(glycolic acid) or poly(ortho esters).

15

A plethora of transdermal patches suitable as delivery systems may be selected from the types described or alluded to in 'Transdermal Drug Delivery; Developmental Issues and Research Initiatives' (Hadgraft and Guy, Marcel Dekker Inc., 1989), by transdermal patches of the type disclosed in US 4,743,249, US 4,906,169, US 5,198,223, US  
 20 4,816,540, or US 5,422,119 or by the transdermal delivery process disclosed in US 5,023,084 or by any transdermal patch known to the person skilled in the art. Moreover, absorption enhancers or skin permeation enhancers of the type disclosed in US 4,379,454 or US 4,973,468 or using any such enhancers known to the person skilled in the art. Transscrotal patches are attractive embodiments of the method of administration.  
 25 Transdermal therapeutic systems, containing the progestin, the androgen, or both components may be based on iontophoresis, diffusion, or a combination of these two effects.

Delivery systems wherein the formulation is administered topically on the skin for  
 30 percutaneous absorption in dosage forms or formulations containing conventionally non-toxic pharmaceutical acceptable carriers and excipients including microspheres and liposomes. Although transdermal patches are one of the preferred methods of percutaneous absorption, it is anticipated that the formulations may be adapted to be suitable for use as creams, ointments, lotions, liniments, gels, hydrogels, solutions,  
 35 suspensions, sticks, sprays, pastes, plasters and other kinds of transdermal drug delivery

systems. The pharmaceutically acceptable carriers or excipients may include emulsifying agents, antioxidants, buffering agents, preservatives, humectants, penetration enhancers, chelating agents, gel-forming agents, ointment bases, perfumes and skin protective agents.

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Examples of emulsifying agents are naturally occurring gums, e.g. gum acacia or gum tragacanth, naturally occurring phosphatides, e.g. soybean lecithin and sorbitan monooleate derivatives.

- 10 Examples of antioxidants are butylated hydroxy anisole (BHA), ascorbic acid and derivatives thereof, tocopherol and derivatives thereof, butylated hydroxy anisole and cysteine.

- Examples of preservatives are parabens, such as methyl or propyl p-hydroxybenzoate  
15 and benzalkonium chloride.

Examples of humectants are glycerin, propylene glycol, sorbitol and urea.

- Examples of penetration enhancers are propylene glycol, DMSO, triethanolamine, *N,N*-  
20 dimethylacetamide, *N,N*-dimethylformamide, 2-pyrrolidone and derivatives thereof, tetrahydrofurfuryl alcohol, propylene glycol, diethylene glycol monoethyl or monomethyl ether with propylene glycol monolaurate or methyl laurate, eucalyptol, lecithin, Transcutol®, and Azone®.

- 25 Examples of chelating agents are sodium EDTA, citric acid and phosphoric acid.

Examples of gel forming agents are Carbopol, cellulose derivatives, bentonite, alginates, gelatin and polyvinylpyrrolidone.

- 30 Subcutaneous implants are well known to the person skilled in the art and a plurality of such implants are suitable methods of administration. Subcutaneous implantation methods are preferably non-irritating and mechanically resilient. The implants may be of matrix type, of reservoir type or hybrids thereof. In matrix type devices, the carrier material may be porous or non-porous, solid or semi-solid, and permeable or  
35 impermeable to the active compound or compounds. The carrier material may be

biodegradable or may slowly erode after administration. In some instances, the matrix is non-degradable but instead relies on the diffusion of the active compound through the matrix for the carrier material to degrade. An alternative subcutaneous implant methods utilises reservoirs devices wherein the active compound or compounds are surrounded by

5 a rate controlling membrane, ideally a membrane independent of component concentration (possessing zero-order kinetics). Devices consisting of a matrix surrounded by a rate controlling membrane are hybrids also anticipated as a method of administration and may optionally be mechanically operated.

- 10 Both reservoir and matrix type devices may comprise of materials such as polydimethylsiloxane, such as Silastic™, or other silicone rubbers. Matrix materials may be insoluble polypropylene, polyethylene, polyvinyl chloride, ethylvinyl acetate, polystyrene and polymethacrylate, as well as glycerol esters of the glycerol palmitostearate, glycerol stearate and glycerol behenate type. Materials may be
- 15 hydrophobic or hydrophilic polymers and optionally comprise of solubilising agents.

In general, subcutaneous implant devices may be slow-release capsules made with any suitable polymer, as described in US 5,035,891 and US 4,210,644, conferring such properties are anticipated and may be of the sort known to the person skilled in the art.

- 20 In general, at least four different approaches are applicable in order to provide rate control over the release and transdermal permeation of a drug compound. These approaches are: membrane-moderated systems, adhesive diffusion-controlled systems, matrix dispersion-type systems and microreservoir systems. It is appreciated that a controlled
- 25 release percutaneous and/or topical composition may be obtained by using a suitable mixture of the above-mentioned approaches.

- In a membrane-moderated system, the active drug substance is present in a reservoir which is totally encapsulated in a shallow compartment molded from a drug-impermeable
- 30 laminate, such as a metallic plastic laminate, and a rate-controlling polymeric membrane such as a microporous or a non-porous polymeric membrane, e.g., ethylene-vinyl acetate copolymer. The active drug substance is only permitted to be released through the ratecontrolling polymeric membrane. In the drug reservoir, the active drug substance may either be dispersed in a solid polymer matrix or suspended in an unleachable, viscous
- 35 liquid medium such as silicone fluid. On the external surface of the polymeric membrane,

a thin layer of an adhesive polymer is applied to achieve an intimate contact of the transdermal system with the skin surface. The adhesive polymer is preferably a polymer which is hypoallergenic and compatible with the active drug substance.

- 5 In an adhesive diffusion-controlled system, a reservoir of the active drug substance is formed by directly dispersing the active drug substance in an adhesive polymer and then - by, e.g., solvent casting - spreading the adhesive containing the active drug substance onto a flat sheet of substantially drug-impermeable metallic plastic backing to form a thin drug reservoir layer.

10

A matrix dispersion-type system is characterized in that a reservoir of the active drug substance is formed by substantially homogeneously dispersing the active drug substance in a hydrophilic or lipophilic polymer matrix and then, the drug-containing polymer is molded into disc with a substantially well-defined surface area and controlled  
15 thickness. The adhesive polymer is spread along the circumference to form a strip of adhesive around the disc.

A microreservoir system may be considered as a combination of the reservoir and matrix dispersion type systems. In this case, the reservoir of the active substance is formed by  
20 first suspending the drug solids in an aqueous solution of water-soluble polymer and then dispersing the drug suspension in a lipophilic polymer to form a multiplicity of unleachable, microscopic spheres of drug reservoirs.

- In embodiments wherein the formulation comprises progestin, non-limiting examples of  
25 methods of administration are outlined above. Moreover, it is anticipated that in embodiments wherein an androgen is further administered, that is to say a combination of two active component substances, it is anticipated that each component may be in separate vials or vesicles or the like. For example, the components may use non-identical vehicles, solvents, buffers, parenterally acceptable carriers and/or excipients or the active  
30 drug substance may independently be incorporated into microspheres, microcapsules, nanoparticles, liposomes or the like for differential controlled release. Furthermore, the compositions may, in addition, conveniently comprise non-identical suspending, solubilising, stabilising, pH-adjusting agents and/or dispersing agents. The non-identical compositions comprise a formulation in one package.

35



In one embodiment of the invention, the formulation comprising both progestin and androgen is administered by non-oral means, such as methods of administrations selected from the group consisting of intramuscular injection, intravenous injection, subcutaneous implantation, subcutaneous injection and transdermal preparations.

5

In an alternative adaptation of the formulation, the methods of administrations of the progestin are selected from the group consisting of intramuscular injection, intravenous injection, subcutaneous implantation, subcutaneous injection and transdermal preparations and the methods of administrations of the androgen are selected from the group consisting of oral administration, intramuscular injection, intravenous injection, subcutaneous implantation, subcutaneous injection and transdermal preparations. That is to say that each active component in the formulation is administered in two separate dosage forms, in two non-identical compositions and/or by two different methods of administration.

15

The preferred methods of administration are those wherein both compounds are administered by a method selected from the group comprising subcutaneous implant, intramuscular injection and transdermal patch.

20 A further aspect of the invention relates to the use of NET, or derivatives thereof, for the preparation of a pharmaceutical composition for use as a male contraceptive. Preferably, the NET derivative is a carboxylic ester, particularly NET enanthate or NET acetate, preferably NET enanthate.

25 Moreover, the invention encompasses to the use of a combination of NET or derivatives thereof and an androgen for the preparation of a male contraceptive formulation. Thus, the use of a combination of NET, or derivatives thereof, and an androgen for the preparation of a preparation of a pharmaceutical composition for use as a male contraceptive is defined by the present invention. In particular, one embodiment of the invention  
30 encompasses the use of a combination of NET esters and testosterone, or derivatives thereof, for the preparation of a pharmaceutical composition for use as a male contraceptive. Preferably, the NET derivative is a carboxylic ester, particularly NET enanthate or NET acetate, preferably NET enanthate. Preferably, the testosterone derivative used is a testosterone ester, particularly testosterone propionate, testosterone

undecanoate, testosterone decanoate, testosterone enanthate, testosterone cypionate and testosterone buciclate, preferably testosterone undecanoate.

It is most preferred that NET enanthate and testosterone undecanoate are used for the  
5 preparation of a pharmaceutical composition for use as male contraceptive.

The invention further encompasses a method of providing male contraception comprising administering to an individual NET or a derivative thereof in an amount sufficient to suppress spermatogenesis such as by administering a non-oral formulation as described  
10 *supra*.

Furthermore, it is preferred that a method of providing male contraception comprises administering to an individual a combination of NET or a derivative thereof and an androgen as a non-oral formulation in an amount sufficient to suppress spermatogenesis.  
15

It is preferred that in such a method, the androgen is testosterone or a derivative thereof, such as a testosterone ester, particularly testosterone undecanoate.

A preferred embodiment of the method of providing male contraception comprises  
20 administering to an individual a combination of NET enanthate and testosterone undecanoate as a non-oral formulation in an amount sufficient to suppress spermatogenesis, particularly by means of a transdermal patch, intramuscular injection or by a subcutaneous implant.

25 The present invention is further described in the following examples which are not in any way intended to limit the scope of the invention as claimed.

## EXAMPLES

### Example 1

#### 5 Clinical Study Design

In this study design (Table1), the three groups A, B and C follow separate regimens. To Group A is administered 200 mg of NET enanthate intramuscularly every six weeks starting day 0 and testosterone undecanoate intramuscularly (1000 mg) on weeks 2 and  
 10 six and every following six weeks. This regimen is followed until week 18. To group B is administered 400 mg of NET enanthate intramuscularly and testosterone undecanoate intramuscularly (1000 mg) every six weeks starting day 0. This regimen was followed until week 18. Group C is administered 10 mg of NET acetate daily *p.o* and 1000 mg of testosterone undecanoate intramuscularly every six weeks for 18 weeks.

15

Examinations and analyses are performed on group members throughout the administration period to establish the relative and absolute efficacy of each of the regimens. Moreover, examinations and analyses are performed until week 56.

20 In studies combining NET and an androgen, the use of TU in trials of male contraception offers considerable advantages compared to other testosterone esters because of its favourable injection interval of 6 weeks. However, the use of other testosterone esters in comparable doses with other injection intervals or by other means of administration offer comparable contraceptive means when used in combination with NET esters. The basis  
 25 for the advantages of the present invention resides in the use of NET esters, alone or in combination with an androgen, preferably by intramuscular or oral administration.

## 1.1.1 Study flow chart

Table 1

			Screening		Start	Treatment										Observation				
Visit	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16				
Assessment	A	B <sup>1</sup>	day 0 <sup>2</sup>	W/2 <sup>3</sup>	W/4	W/6	W/8	W/12	W/16	W/18	W/20	W/24	W/28	W/36	W/44	W/56				
Group A																				
NET-EY administration (mg)			200			200		200		200										
TU administration (mg)				1000		1000		1000		1000										
Group B																				
NET-EY administration (mg)			400			400		400		400										
TU administration (mg)			1000			1000		1000		1000										
Group C																				
NET-A administration (mg)			10 mg																	
Concomitant of visit			V 1/2																	
TU administration (mg)			1000																	
Informed Consent/Denial/refusals	ABC																			
Check of Inclusion/Exclusion	ABC		ABC																	
Randomization/Urinary dispensed			ABC																	
Medical/and/or medication history	ABC	ABC	ABC																	
Vital signs	ABC	ABC				ABC		ABC		ABC		ABC		ABC		ABC				
Physical examination	ABC					ABC		ABC		ABC		ABC		ABC		ABC				
Andrological examination	ABC					ABC		ABC		ABC		ABC		ABC		ABC				
Semen analysis	ABC	ABC				ABC		ABC		ABC		ABC		ABC		ABC				
Genital laboratory	ABC							ABC				ABC				ABC				
Hormones	ABC	ABC		A		ABC	ABC	ABC	ABC	ABC	ABC	ABC	ABC	ABC	ABC	ABC				
Exam/Ultrasoundography of prostate		ABC						ABC				ABC			ABC	ABC				
Ultrasonography of testes		ABC						ABC				ABC			ABC	ABC				
Return of diagnostics, if available																ABC				
Anti-concomitant medication				A		ABC	ABC	ABC	ABC	ABC	ABC	ABC	ABC	ABC	ABC	ABC				
Questionnaire	ABC	ABC			ABC		ABC		ABC		ABC		ABC		ABC	ABC				
At least 7 days after visit 1																				
Not more than 8 weeks after visit 1																				
A deviation of + 5 days will be tolerated																				

<sup>1</sup> At least 7 days after visit 1<sup>2</sup> Not more than 8 weeks after visit 1<sup>3</sup> A deviation of  $\pm 5$  days will be tolerated

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## Example 2

### Estrogenic Properties

- 5 Receptor binding assays, transactivation assays, and *in vivo* assays for establishing estrogenic activity are described in *Handbook of experimental Pharmacology, Vol. 135/II, Estrogens and Antiestrogens II, Pharmacology and Clinical Applications of Estrogens and Antiestrogens*; M. Ottel and E. Schillinger (editors); K.-H. Fritzemeier and C. Hegele-Hartung, *In Vitro and In Vivo Models to Characterise Estrogens and Antiestrogens*;  
10 Springer-Verlag, Berlin, Heidelberg, 1999.

The basic principle behind receptor binding assays lies in that the binding of substances to the estrogen receptor is a pre-requisite for the compound to exhibit estrogen-like activity. Binding affinities or dissociation constants are used as measures of the  
15 substance to bind to the estrogen receptor.

Transactivation assays for the detection of estrogenic activity are based on the ability of the estrogen receptor to cause gene activation in a ligand-dependant way. The binding of an estrogenic substance results in ligand-activated formation of a receptor dimer which  
20 then binds to a specific nucleotide sequence in the promoter segment of a target gene.

A number a of test systems and method for the *in vivo* characterisation of estrogenic activity. The Allen-Doisy Assay is one such assay in which the vaginal cytology of ovariectomised rodents is examined. Given estrogenic compounds induce proliferation  
25 and keratinisation of the vaginal epithelium, the microscopic examination of vaginal smears of ovariectomised rodents for evaluation of epithelial thickness and cornification is a reliable indicator of estrogenic activity.

Other *in vivo* test systems to test estrogenicity in the vagina include determination of  
30 alterations in the vaginal epithelial mitotic index, of a reduction in vaginal tetrazolium, of the vaginal opening in immature rodents and of the vaginatrophic response as well as by the measurement of sialic acid production in ovariectomised female rodents.

Many assays consisting of determining the estrogenicity in the uterus of rodents and  
35 primates are known to the person skilled in the art.

Tests on the influence of estrogens on hypothalamic-pituitary-ovarian feedback typically comprises of determining the change in gonadal weight given the principle that the administration of gonadal steroids leads of reduced growth of the ovaries and testes.

- 5 Alternatively, the reduction in the peak levels of the gonadotropins FSH/LH during the preovulatory phase in female rodents can be a measured.

### Example 3

#### 10 Androgenic Properties

Assays, both *in vivo* and *in vitro*, for establishing androgenic activity are described in *Androgens II and Anti-Androgens*, F. Neuman, F. Baher, J. Brotherton, K.-J. Gräf, S.H. Hasan, H.J. Horn, A. Hughes, G.W. Oertel, H. Steinbeck, H.E. Voss, R.K. Wagner,

- 15 Springer-Verlag, Berlin, Heidelberg, 1974.

Receptor tests for androgenicity are analogous to estrogenicity tests and generally utilise rat prostate cytosol for tests on androgen receptors. A transfection assay of the androgen receptor has been established by Fuhrmann, Bengston, Repenthin, and Schillinger (*J.*

- 20 *Steroid Biochem. Mol. Biol.*, 1992, 42(8), 787). An antiproliferation test with the human prostate cancer cell line LCNaP, which expresses the androgen receptor and can be stimulated in growth by androgens has been developed. If a single administration of a test compound leads to growth stimulation, this can be explained by the androgenic activity of the compound.

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